

### **AMENDMENTS TO THE CLAIMS**

This listing of the claims will replace all prior versions, and listings, of the claims in the application.

#### **Listing of Claims**

1. (Currently Amended) A method for treating a mammalian subject having a solid tumor *ex vivo*, comprising, direct injection of a nucleic acid molecule encoding:

a) at least 20 contiguous amino acids of SEQ ID NO: 2 or 4; or

b) a polypeptide that is at least 50% homologous to SEQ ID NO: 2 or 4;

into cells of the tumor, wherein said nucleic acid molecule encodes encoding a B7-2 molecule or a fragment thereof in a form suitable for expression of the B7-2 molecule, into cells of the tumor, and wherein the B7-2 molecule or fragment thereof has the ability to costimulate a T cell and the ability to bind a CD28 or CTLA4 ligand, such that the growth of the tumor is inhibited.

2. (Currently Amended) A method for modifying cells of a solid tumor *ex vivo* to express a B7-2 molecule or a fragment thereof comprising, direct injection of a nucleic acid molecule encoding:

a) at least 20 contiguous amino acids of SEQ ID NO: 2 or 4; or

b) a polypeptide that is at least 50% homologous to SEQ ID NO: 2 or 4;

into cells of the tumor, wherein said nucleic acid molecule encodes encoding a B7-2 molecule or a fragment thereof in a form suitable for expression of the B7-2 molecule, into cells of the tumor, and wherein the B7-2 molecule or fragment thereof has the ability to costimulate a T cell and the ability to bind a CD28 or CTLA4 ligand, such that B7-2 is expressed by the tumor cells.

3. (Currently Amended) A method of increasing the immunogenicity immunogenicity of a cells of a solid tumor *ex vivo* comprising, direct injection of a nucleic acid molecule encoding:

a) at least 20 contiguous amino acids of SEQ ID NO: 2 or 4; or

b) a polypeptide that is at least 50% homologous to SEQ ID NO: 2 or 4;  
into cells of the tumor, wherein said nucleic acid molecule encodes encoding a B7-2 molecule or a fragment thereof in a form suitable for expression of the B7-2 molecule, into cells of the tumor, and wherein the B7-2 molecule or fragment thereof has the ability to costimulate a T cell and the ability to bind a CD28 or CTLA4 ligand, such that B7-2 is expressed by the tumor cells, to thereby increasing increase the immunogenicity of the tumor cells.

4. (Currently Amended) The method of any of claims 1-3, wherein the nucleic acid molecule encoding a B7-2 molecule comprises the nucleic acid sequence shown in SEQ ID NO: 1 or SEQ ID NO: 3.

5. (Currently Amended) The method of any of claims 1-3, wherein B7-2 comprises the amino acid sequence shown in SEQ ID NO:2 or SEQ ID NO: 4.

6. (Original) The method of any of claims 1-3, wherein the nucleic acid molecule encoding B7-2 is in a viral vector.

7. (Original) The method of claim 6, wherein the viral vector is selected from the group consisting of a retroviral vector, an adenoviral vector, and an adeno-associated viral vector.

8. (Original) The method of any of claims 1-3, wherein the nucleic acid molecule encoding B7-2 is a plasmid expression vector.

9. (Original) The method of any of claims 1-3, wherein the tumor cells are further transfected with at least one nucleic acid molecule encoding a B7-3 protein.

10. (Original) The method of any of claims 1-3, wherein the tumor cells are further injected with at least one nucleic acid molecule encoding at least one MHC class II  $\alpha$  chain protein and at least one MHC class II  $\beta$  chain protein in a form suitable for expression of the MHC class II  $\alpha$  chain protein(s) and the MHC class II  $\beta$  chain protein(s).

11. (Original) The method of any of claims 1-3, wherein the tumor cells are further injected with at least one nucleic acid molecule encoding at least one MHC class I  $\alpha$  chain protein in a form suitable for expression of the MHC class I protein(s).

12. (Original) The method of any of claims 1-3, wherein the tumor cells are further injected with a nucleic acid molecule encoding a  $\beta$ -2 microglobulin protein in a form suitable for expression of the  $\beta$ -2 microglobulin protein.

13. (Original) The method of any of claims 1-3, wherein expression of the MHC class II invariant chain is inhibited in the tumor cells by transfection of the tumor cells with a nucleic acid molecule which is antisense to a regulatory or a coding region of the invariant chain gene.

14. (Original) The method of any of claims 1-3 wherein the solid tumor is selected from a group consisting of a carcinoma, sarcoma, melanoma and neuroblastoma.